to assist the implementation of new programs and national strategies for dissemination of information intended to prevent alcohol and drug abuse:

- (3) encourage media outlets throughout the country to provide information aimed at preventing alcohol and drug abuse, including public service announcements, documentary films, and advertisements; and
- (4) evaluate the effectiveness and assist in the update of programs and national strategies formulated with the assistance of the Commission

(Pub. L. 99–570, title VIII, §8003, Oct. 27, 1986, 100 Stat. 3207–161.)

### § 1303. Membership

### (a) Number and appointment

The Commission shall be composed of 12 members appointed by the President within 30 days after October 27, 1986, and should include representatives of—

- (1) advertising agencies;
- (2) motion picture, television, radio, cable communications, and print media;
  - (3) the recording industry;
- (4) other segments of the business sector of the United States;
- (5) experts in the prevention of alcohol and drug abuse;
- (6) professional sports organizations and associations; and
- (7) other Federal agencies, as designated by the President, including the Director of the Agency for Substance Abuse Prevention of the Department of Health and Human Services.

### (b) Terms

- (1) Except as provided in paragraphs (2) and (3), members shall be appointed for terms of 3 years.
- (2) Any member appointed to fill a vacancy occurring before the expiration of the term for which his predecessor was appointed shall be appointed only for the remainder of such term.
- (3) A member may serve after the expiration of his term until his successor has taken office.

# (c) Basic pay and expenses

- (1) Except as provided in paragraph (2), members of the Commission shall serve without pay.
- (2) While away from their homes or regular places of business in the performance of services for the Commission, members shall be allowed travel expenses, including a per diem allowance in lieu of subsistence, in the same manner as persons serving intermittently in the Government service are allowed travel expenses under section 5703 of title 5.

(Pub. L. 99–570, title VIII, §8004, Oct. 27, 1986, 100 Stat. 3207–162.)

SECTION REFERRED TO IN OTHER SECTIONS

This section is referred to in section 1304 of this title.

## § 1304. Meetings

## (a) In general

- (1) The Commission shall meet at the call of the Moderator.
- (2) The Moderator shall convene the 1st meeting of the Commission within 30 days after the

date of the completion of appointments under section 1303(a) of this title.

### (b) Moderator

One member of the Commission shall be designated by the President to serve as Moderator of the Commission.

### (c) Quorum and procedure

The Commission shall adopt rules regarding quorum requirements and meeting procedures as the Commission deems appropriate at the 1st meeting of the Commission.

### (d) Voting

Decisions and official acts of the Commission shall be according to the vote of a majority of members at a properly called meeting.

(Pub. L. 99–570, title VIII, §8005, Oct. 27, 1986, 100 Stat. 3207–162.)

# § 1305. Director and staff; experts and consultants

#### (a) Director and staff

- (1) Subject to paragraph (2), the Moderator, with the approval of the Commission, may employ and set the rate of pay for a Director and such staff as the Moderator deems necessary.
- (2) Rates of pay set under paragraph (1) shall be less than the rate of basic pay payable under section 5316 of title 5.

### (b) Experts and consultants

The Moderator, with the approval of the Commission, may procure temporary and intermittent services under section 3109(b) of title 5.

### (c) Staff of Federal agencies

Upon request of the Commission, the head of any Federal agency is authorized to detail, on a reimbursable basis, any of the personnel of such agency to the Commission to assist the Commission in carrying out its duties under this chapter.

(Pub. L. 99–570, title VIII, §8006, Oct. 27, 1986, 100 Stat. 3207–163.)

## § 1306. Powers of Commission

## (a) Hearings and sessions

The Commission may, for the purpose of carrying out this chapter, hold such hearings, sit and act at such times and places, take such testimony, and receive such evidence, as the Commission considers appropriate.

### (b) Obtaining official data

Upon the request of the Moderator of the Commission, the Commission may secure directly from any department or agency of the United States information necessary to enable it to carry out this chapter.

### (c) Gifts

The Commission may accept, use, and dispose of gifts or donations of services or property.

### (d) Mails

The Commission may use the United States mails in the same manner and under the same conditions as other departments and agencies of the United States.

## (e) Administrative support services

The Administrator of General Services shall provide to the Commission on a reimbursable basis such administrative support services as the Commission may request.

(Pub. L. 99–570, title VIII, §8007, Oct. 27, 1986, 100 Stat. 3207–163.)

## § 1307. Report

The Commission shall transmit to the President and to each House of Congress a report not later than July 31 of each year which contains a detailed statement of the activities of the Commission during the preceding year, including a summary of the number of public service announcements produced by the Commission and published or broadcast.

(Pub. L. 99–570, title VIII, \$8008, Oct. 27, 1986, 100 Stat. 3207-163.)

TERMINATION OF REPORTING REQUIREMENTS

For termination, effective May 15, 2000, of provisions in this section relating to transmitting an annual report to each House of Congress, see section 3003 of Pub. L. 104-66, as amended, set out as a note under section 1113 of Title 31, Money and Finance, and page 18 of House Document No. 103-7.

#### § 1308. Termination

The Commission shall terminate on a date which is three years after the date on which members of the Commission are first appointed, unless the President, by Executive order, extends the authority of the Commission.

(Pub. L. 99–570, title VIII, §8009, Oct. 27, 1986, 100 Stat. 3207–163.)

# CHAPTER 19—PESTICIDE MONITORING IMPROVEMENTS

Sec.

1401. Pesticide monitoring and enforcement information.

- (a) Data management systems.
- (b) Information.
- (c) Volume data.
- (d) Compilation.

1402. Foreign pesticide information.

- (a) Cooperative agreements.
- (b) Information activities.
- (c) Coordination with other agencies.
- (d) Report.

1403. Pesticide analytical methods.

# § 1401. Pesticide monitoring and enforcement information

# (a) Data management systems

- (1) Not later than 480 days after August 23, 1988, the Secretary of Health and Human Services shall place in effect computerized data management systems for the Food and Drug Administration under which the Administration will—
  - (A) record, summarize, and evaluate the results of its program for monitoring food products for pesticide residues,
  - (B) identify gaps in its pesticide monitoring program in the monitoring of (i) pesticides, (ii) food products, and (iii) food from specific countries and from domestic sources,
  - (C) detect trends in the presence of pesticide residues in food products and identify public

health problems emerging from the occurrence of pesticide residues in food products,

- (D) focus its testing resources for monitoring pesticide residues in food on detecting those residues which pose a public health concern,
- (E) prepare summaries of the information listed in subsection (b) of this section, and
- (F) provide information to assist the Environmental Protection Agency in carrying out its responsibilities under the Federal Insecticide, Fungicide, and Rodenticide Act [7 U.S.C. 136 et seq.] and the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 301 et seq.].
- (2) As soon as practicable, the Secretary of Health and Human Services shall develop a means to enable the computerized data management systems placed into effect under paragraph (1) to make the summary described in subsection (c) of this section.
- (3)(A) Paragraph (1) does not limit the authority of the Food and Drug Administration to—
- (i) use the computerized data management systems placed in effect under paragraph (1), or
- (ii) develop additional data management systems.

to facilitate the regulation of any substance or product covered under the requirements of the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 301 et seq.].

(B) In placing into effect the computerized data management systems under paragraph (1) and in carrying out paragraph (2), the Secretary shall comply with applicable regulations governing computer system design and procurement.

## (b) Information

The Food and Drug Administration shall use the computerized data management systems placed into effect under subsection (a)(1) of this section to prepare a summary of—

- (1) information on-
- (A) the types of imported and domestically produced food products analyzed for compliance with the requirements of the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 301 et seq.] regarding the presence of pesticide residues,
- (B) the number of samples of each such food product analyzed for such compliance by country of origin,
- (C) the pesticide residues which may be detected using the testing methods employed,
- (D) the pesticide residues in such food detected and the levels detected,
- (E) the compliance status of each sample of such food tested and the violation rate for each country-product combination, and
- (F) the action taken with respect to each sample of such food found to be in violation of the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 301 et seq.] and its ultimate disposition, and
- (2) information on—
- (A) the country of origin of each imported food product referred to in paragraph (1)(A), and
- (B) the United States district of entry for each such imported food product.

TITLE 21—FOOD AND DRUGS

## (c) Volume data

The Food and Drug Administration shall use the computerized data management systems placed into effect under subsection (a)(1) of this section to summarize the volume of each type of food product subject to the requirements of the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 301 et seq.] which is imported into the United States and which has an entry value which exceeds an amount established by the Secretary of Health and Human Services. The summary shall be made by country of origin and district of entry. Information with respect to volumes of food products to be included in the summary shall, to the extent feasible, be obtained from data bases of other Federal agencies.

## (d) Compilation

Not later than 90 days after the expiration of 1 year after the data management systems are placed into effect under subsection (a) of this section and annually thereafter, the Secretary of Health and Human Services shall compile a summary of the information described in subsection (b) of this section with respect to the previous year. When the Food and Drug Administration is able to make summaries under subsection (c) of this section, the Secretary shall include in the compilation under the preceding sentence a compilation of the information described in subsection (c) of this section. Compilations under this subsection shall be made available to Federal and State agencies and other interested persons.

(Pub. L. 100–418, title IV, §4702, Aug. 23, 1988, 102 Stat. 1412.)

### REFERENCES IN TEXT

The Federal Insecticide, Fungicide, and Rodenticide Act, referred to in subsec. (a)(1)(F), is act June 25, 1947, ch. 125, as amended generally by Pub. L. 92–516, Oct. 21, 1972, 86 Stat. 973, which is classified generally to subchapter II (§136 et seq.) of chapter 6 of Title 7, Agriculture. For complete classification of this Act to the Code, see Short Title note set out under section 136 of Title 7 and Tables.

The Federal Food, Drug, and Cosmetic Act, referred to in subsecs. (a) to (c), is act June 25, 1938, ch. 675, 52 Stat. 1040, as amended, which is classified generally to chapter 9 (§301 et seq.) of this title. For complete classification of this Act to the Code, see section 301 of this title and Tables.

### SHORT TITLE

Section 4701 of Pub. L. 100–418 provided that: "This subtitle [subtitle G (§§ 4701–4704) of title IV of Pub. L. 100–418, enacting this chapter] may be cited as the 'Pesticide Monitoring Improvements Act of 1988'."

IMPORTED MEAT, POULTRY PRODUCTS, EGGS, AND EGG PRODUCTS

Section 4506 of Pub. L. 100-418 provided that:

"(a) REPORT.—Not later than 90 days after the date of the enactment of this Act [Aug. 23, 1988], the Secretary of Agriculture shall submit a report to Congress— "(1) specifying the planned distribution, in fiscal

"(1) specifying the planned distribution, in fiscal years 1988 and 1989, of the resources of the Department of Agriculture available for sampling imported covered products to ensure compliance with the requirements of the Federal Meat Inspection Act (21 U.S.C. 601 et seq.), the Poultry Products Inspection Act (21 U.S.C. 451 et seq.), and the Egg Products Inspection Act (21 U.S.C. 1031 et seq.) that govern the level of residues of pesticides, drugs, and other products permitted in or on such products;

"(2) describing current methods used by the Secretary to enforce the requirements of such Acts with respect to the level of residues of pesticides, drugs, and other products permitted in or on such products;

"(3) responding to the audit report of the Inspector General of the Department of Agriculture, Number 38002—2—hy, dated January 14, 1987;

"(4) providing a summary with respect to the importation of covered products during fiscal years 1987 and 1988 that specifies—

"(A) the number of samples of each such product taken during each such fiscal year in carrying out the requirements described in paragraph (1); and

"(B) for each violation of such requirements during each such fiscal year—

"(i) the covered products with respect to which such violation occurred;

"(ii) the residue in or on such product in violation of such requirements;

"(iii) the country exporting such product;

"(iv) the actions taken in response to such violation and the reasons for such actions; and

"(v) the level of testing conducted by the countries exporting such products;

"(5) describing any research conducted by the Secretary to develop improved methods to detect residues subject to such requirements in or on covered products; and

"(6) providing any recommendations the Secretary considers appropriate for legislation to add or modify penalties for violations of laws, regulations, and other enforcement requirements governing the level of residues that are permitted in or on imported covered products.

"(b) REVISION.—Not later than November 15, 1989, the Secretary of Agriculture shall revise, as necessary, the report prepared under subsection (a) and submit the revision to Congress.

"(c) DEFINITION.—As used in this section, the term 'covered products' means meat, poultry products, eggs, and egg products."

## § 1402. Foreign pesticide information

### (a) Cooperative agreements

The Secretary of Health and Human Services shall enter into cooperative agreements with the governments of the countries which are the major sources of food imports into the United States subject to pesticide residue monitoring by the Food and Drug Administration for the purpose of improving the ability of the Food and Drug Administration to assure compliance with the pesticide tolerance requirements of the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 301 et seq.] with regard to imported food.

## (b) Information activities

(1) The cooperative agreements entered into under subsection (a) of this section with governments of foreign countries shall specify the action to be taken by the parties to the agreements to accomplish the purpose described in subsection (a) of this section, including the means by which the governments of the foreign countries will provide to the Secretary of Health and Human Services current information identifying each of the pesticides used in the production, transportation, and storage of food products imported from production regions of such countries into the United States.

(2) In the case of a foreign country with which the Secretary is unable to enter into an agreement under subsection (a) of this section or for which the information provided under paragraph (1) is insufficient to assure an effective pesticide monitoring program, the Secretary shall, to the extent practicable, obtain the information described in paragraph (1) with respect to such country from other Federal or international agencies or private sources.

- (3) The Secretary of Health and Human Services shall assure that appropriate offices of the Food and Drug Administration which are engaged in the monitoring of imported food for pesticide residues receive the information obtained under paragraph (1) or (2).
- (4) The Secretary of Health and Human Services shall make available any information obtained under paragraph (1) or (2) to State agencies engaged in the monitoring of imported food for pesticide residues other than information obtained from private sources the disclosure of which to such agencies is restricted.

### (c) Coordination with other agencies

The Secretary of Health and Human Services shall—

- (1) notify in writing the Department of Agriculture, the Environmental Protection Agency, and the Department of State at the initiation of negotiations with a foreign country to develop a cooperative agreement under subsection (a) of this section; and
- (2) coordinate the activities of the Department of Health and Human Services with the activities of those departments and agencies, as appropriate, during the course of such negotiations.

### (d) Report

Not later than one year after August 23, 1988, the Secretary of Health and Human Services shall report to the Committee on Agriculture, Nutrition, and Forestry and the Committee on Labor and Human Resources of the Senate and the House of Representatives on the activities undertaken by the Secretary to implement this section. The report shall be made available to appropriate Federal and State agencies and to interested persons.

(Pub. L. 100–418, title IV,  $\S4703$ , Aug. 23, 1988, 102 Stat. 1413.)

### References in Text

The Federal Food, Drug, and Cosmetic Act, referred to in subsec. (a), is act June 25, 1938, ch. 675, 52 Stat. 1040, as amended, which is classified generally to chapter 9 (§301 et seq.) of this title. For complete classification of this Act to the Code, see section 301 of this title and Tables.

## § 1403. Pesticide analytical methods

The Secretary of Health and Human Services shall, in consultation with the Administrator of the Environmental Protection Agency—

- (1) develop a detailed long-range plan and timetable for research that is necessary for the development of and validation of—
  - (A) new and improved analytical methods capable of detecting at one time the presence of multiple pesticide residues in food, and
  - (B) rapid pesticide analytical methods, and
- (2) conduct a review to determine whether the use of rapid pesticide analytical methods by the Secretary would enable the Secretary

to improve the cost-effectiveness of monitoring and enforcement activities under the Federal Food, Drug, and Cosmetic Act [21 U.S.C. 301 et seq.], including increasing the number of pesticide residues which can be detected and the number of tests for pesticide residues which can be conducted in a cost-effective manner.

The Secretary shall report the plan developed under paragraph (1), the resources necessary to carry out the research described in such paragraph, recommendations for the implementation of such research, and the result of the review conducted under paragraph (2) not later than the expiration of 240 days after August 23, 1988, to the Committee on Agriculture, Nutrition, and Forestry and the Committee on Labor and Human Resources of the Senate and the House of Representatives.

(Pub. L. 100–418, title IV, §4704, Aug. 23, 1988, 102 Stat. 1414.)

### REFERENCES IN TEXT

The Federal Food, Drug, and Cosmetic Act, referred to in text, is act June 25, 1938, ch. 675, 52 Stat. 1040, as amended, which is classified generally to chapter 9 (§301 et seq.) of this title. For complete classification of this Act to the Code, see section 301 of this title and Tables

# CHAPTER 20—NATIONAL DRUG CONTROL PROGRAM

SUBCHAPTER I—OFFICE OF NATIONAL DRUG CONTROL POLICY

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